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NEWS	4	JAN 28	USPATFULL, USPAT2, and USPATOLD enhanced with new custom IPC display formats
NEWS	5	JAN 28	MARPAT searching enhanced
NEWS	6	JAN 28	USGENE now provides USPTO sequence data within 3 days of publication
NEWS	7	JAN 28	TOXCENTER enhanced with reloaded MEDLINE segment
NEWS	8	JAN 28	MEDLINE and LMEMLINE reloaded with enhancements
NEWS	9	FEB 08	STN Express, Version 8.3, now available
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NEWS	11	FEB 25	IFIREF reloaded with enhancements
NEWS	12	FEB 25	IMSPRODUCT reloaded with enhancements
NEWS	13	FEB 29	WPINDEX/WPIDS/WPIX enhanced with ECLA and current U.S. National Patent Classification
NEWS	14	MAR 31	IFICDB, IFIPAT, and IFIUDB enhanced with new custom IPC display formats
NEWS	15	MAR 31	CAS REGISTRY enhanced with additional experimental spectra
NEWS	16	MAR 31	CA/CAPLUS and CASREACT patent number format for U.S. applications updated
NEWS	17	MAR 31	LPCI now available as a replacement to LDPCI
NEWS	18	MAR 31	EMBASE, EMBAL, and LEMBASE reloaded with enhancements
NEWS	19	APR 04	STN AnaVist, Version 1, to be discontinued
NEWS EXPRESS	FEBRUARY 08 CURRENT WINDOWS VERSION IS V8.3, AND CURRENT DISCOVER FILE IS DATED 20 FEBRUARY 2008		
NEWS HOURS	STN Operating Hours Plus Help Desk Availability		
NEWS LOGIN	Welcome Banner and News Items		
NEWS IPC8	For general information regarding STN implementation of IPC 8		

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=> S CYHALOTHRIN

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2 CYHALOTHRINS

L1 1726 CYHALOTHRIN

(CYHALOTHRIN OR CYHALOTHRINS)

=> S L1 AND CHLORINATION

81328 CHLORINATION

447 CHLORINATIONS

81401 CHLORINATION

(CHLORINATION OR CHLORINATIONS)

L2 5 L1 AND CHLORINATION

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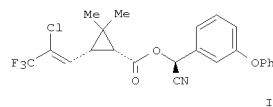
L2 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2006:346943 CAPLUS
DOCUMENT NUMBER: 146:100882
TITLE: Synthesis of carboxylic acid chlorides from
bis(trichloromethyl) carbonate
AUTHOR(S): Xu, Xiangsheng; Du, Xiaohua; Zheng, Mei; Xu, Zhenyuan
CORPORATE SOURCE: Catalytic Hydrogenation Research Center, Zhejiang
University of Technology, Hangzhou, 310014, Peop.
Rep. China
SOURCE: Nongyao (2005), 44(6), 265-266
CODEN: NONGFF; ISSN: 1006-0413
PUBLISHER: Nongyao Bianjibu
DOCUMENT TYPE: Journal
LANGUAGE: Chinese
OTHER SOURCE(S): CASREACT 146:100882

AB Eight representative carboxylic acid chlorides were synthesized by
treatment of the corresponding carboxylic acids with bis
(trichloromethyl)
carbonate. The chloride compds. were cis/trans-3-(2,2-dichloroethenyl)-
2,2-dimethylcyclopropanecarbonyl chloride,
cis-3-(2-chloro-3,3,3-trifluoro-
1-propenyl)-2,2-dimethylcyclopropanecarbonylchloride, chloroacetyl
chloride, trichloroacetyl chloride, benzoyl chloride, 2-chlorobenzoyl
chloride, 2-methylbenzoyl chloride, and trans-3-phenyl-2-propenoyl
chloride. Yields and purity of the products were 82.1% to 91.7% and
96.4%
to 98.9%, resp.

L2 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2004:718504 CAPLUS
DOCUMENT NUMBER: 141:243704
TITLE: Process for preparing gamma-cyhalothrin
INVENTOR(S): Brown, Stephen Martin; Gott, Brian David
PATENT ASSIGNEE(S): Syngenta Limited, UK
SOURCE: PCT Int. Appl., 18 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004074237	A1	20040902	WO 2004-GB726	20040223
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GR, GU, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MY, NA, NI, RW: BW, GH, GM, KE, LS, MM, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
CA 2512429	A1	20040902	CA 2004-2512429	20040223
EP 1599442	A1	20051130	EP 2004-713588	20040223
EP 1599442	B1	20070808		
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK			
BR 2004007774	A	20060214	BR 2004-7774	20040223
CN 1738792	A	20060222	CN 2004-80002246	20040223
JP 2006518729	T	20060817	JP 2006-502318	20040223
AT 369332	T	20070815	AT 2004-713588	20040223
ES 2287702	T3	20071216	ES 2004-713588	20040223
US 20060148892	A1	20060706	US 2005-546138	20050819
IN 2005CN02002	A	20070727	IN 2005-CN2002	20050823
PRIORITY APPLN. INFO.:			GB 2003-4132	A 20030224
			WO 2004-GB726	W 20040223

OTHER SOURCE(S): CASREACT 141:243704
GI



AB A process for the preparation of gamma-cyhalothrin (I) comprising

L2 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN (Continued)
steps of (a) chlorinating (1R)-cis-(Z)-3-(2-chloro-3,3,3-trifluoro-1-propenyl)-2,2-dimethylcyclopropanecarboxylic acid to give
(1R)-cis-(Z)-3-(2-chloro-3,3,3-trifluoro-1-propenyl)-2,2-dimethylcyclopropanecarboxylic acid chloride and (b) esterifying
(1R)-cis-(Z)-3-(2-chloro-3,3,3-trifluoro-1-propenyl)-2,2-dimethylcyclopropanecarboxylic acid chloride with the (S)-cyanohydrin of
3-phenoxybenzaldehyde.
REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE
FORMAT

L2 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2003:511276 CAPLUS
DOCUMENT NUMBER: 139:85510
TITLE: A process for the production of 1R pyrethroid esters
via resolution of cyclopropanecarboxylic acids
Brown, Stephen Martin; Gott, Brian David
INVENTOR(S): Syngenta Limited, UK
PATENT ASSIGNEE(S): Syngenta Limited, UK
SOURCE: PCT Int. Appl., 40 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003053905	A1	20030703	WO 2002-GB5467	20021204
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, GR, GU, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MY, NA, NI, RW: BW, GH, GM, KE, LS, MM, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
AU 2002366752	A1	20030709	AU 2002-366752	20021204
PRIORITY APPLN. INFO.:			GB 2001-30517	A 20011220
			WO 2002-GB5467	W 20021204

OTHER SOURCE(S): CASREACT 139:85510; MARPAT 139:85510
GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

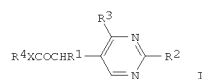
AB A process for the production of 1R pyrethroid esters I [A, B = chlorine
or
bromine or one of A or B is chlorine and the other is trifluoromethyl; R
=
a pyrethroid alc. fragment] or II, which process comprises (a) resolving
pyrethroid acids III where A and B are as defined above to give a
substantially pure 1R cis enantiomer, (b) recovering the 1S cis
enantiomer, (c) optionally converting the 1S cis enantiomer acid to a 1S
cis enantiomer anhydride, acid chloride or pyrethroid ester containing
the
group R where R is a pyrethroid alc. fragment; (d) converting the 1S cis
enantiomer from step b or step c to the 1R trans isomer; (e) optionally
purifying the 1R trans isomer from step d and recycle of the unconverted
1S cis isomer back to step c or d, (f) converting the 1R cis isomer of
the
acid from step a into corresponding 1R cis isomers of the pyrethroid
esters alone, or together with the product of step d or e where the
product of step d or e is not already a pyrethroid ester containing the
group

L2 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN (Continued)
 R. Thus, (1R)-trans-tefluthrin (IV) was prepd. from (±)-cis-(Z)-3-(2-chloro-3,3,3-trifluoro-1-propenyl)-2,2-dimethylcyclopropanecarboxylic acid, via enantiomer resolu. with (R)-(+)-α-methylbenzylamine to give the (1S)-cis-isomer, chlorination with SOCl₂ in the presence of Et₃N, thermal isomerization to the (1R)-trans-acid, chlorination with SOCl₂ and esterification with 2,3,5,6-tetrafluoro-4-methylbenzyl alc. The pesticidal and insecticidal activity of IV was detd. [LC50 = 1.8 & LC90 = 4.6 vs. *Heliothis virescens*; LC50 = 14.285 (resistance factor 35) vs. *Plutella xylostella*].
 REFERENCE COUNT: 14 THERE ARE 14 CITED REFERENCES AVAILABLE FOR THIS
 RECORD. ALL CITATIONS AVAILABLE IN THE RE
 FORMAT

L2 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1989:553826 CAPLUS
 DOCUMENT NUMBER: 111:153826
 ORIGINAL REFERENCE NO.: 111:25660h,25661a
 TITLE: Preparation of pyrimidine-containing carboxylic acid esters having insecticidal and acaricidal activities
 McDonald, Edward; Salmon, Roger; Whittle, Alan John; Hutchings, Michael Gordon
 INVENTOR(S): Imperial Chemical Industries PLC, UK
 PATENT ASSIGNEE(S): Eur. Pat. Appl., 104 pp.
 SOURCE: CODEN: EPXXDW
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 3
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 295839	A2	19881221	EP 1988-305337	19880610
EP 295839	A3	19910731		
R: AT, BE, CH, DE, ES, FR, GB, GR, IT, LI, LU, NL, SE				
ZA 8803862	A	19890222	ZA 1988-3862	19880530
AU 8817389	A	19881222	AU 1988-17389	19880603
AU 610184	B2	19910516		
GB 2209525	A	19890517	GB 1988-13780	19880610
GB 2209525	B	19910403		
HU 47384	A2	19890328	HU 1988-3052	19880615
HU 203644	B	19910930		
BR 8802952	A	19890110	BR 1988-2952	19880616
DK 8803348	A	19881218	DK 1988-3348	19880617
CN 1030412	A	19890118	CN 1988-103836	19880617
CN 1019574	B	19921223		
JP 01016769	A	19890120	JP 1988-148425	19880617
SU 1801108	A3	19930307	SU 1988-4613066	19881212
			GB 1987-14233	A 19870617

PRIORITY APPLN. INFO.:
 OTHER SOURCE(S): CASREACT 111:153826; MARPAT 111:153826
 GI



AB The title compds. [I; R1 = C1-6 alkyl, C2-8 alkenyl, C2-6 alkynyl, C1-4 haloalkyl, C2-8 haloalkenyl, C3-6 cycloalkyl optionally substituted by ≥1 C1-4 alkyl or halo; R2 = C1-8 alkyl, C1-4 haloalkyl, C1-6 alkoxy, halo, C3-6 cycloalkyl optionally substituted by ≥1 C1-4 alkyl or halo, Ph optionally substituted by ≥1 C1-4 alkyl, C1-4 haloalkyl, or C1-4 alkoxy; R3 = H, halo; R4 = residue of an alc. of formula R4-OH which forms an insecticidal ester when combined with

L2 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN (Continued)
 chrysanthemic acid, permethrin, or cyhalothrin acid; X = O, S], useful as insecticides or acaricides, were prepd. To a stirred soln. of 0.1 (RS)-2-[2-[(1,1-dimethylethyl)pyrimidin-5-yl]-3,3-dimethylbutanoic acid, 0.089 2,3,5,6-tetrafluoro-4-(methoxymethyl)benzyl alc., and 0.002 g 4-dimethylaminopyridine in CH₂Cl₂, 0.084 g DCC was added and the mixt. was stirred 18 h to give 0.09 g 2,3,5,6-tetrafluoro-4-(methoxymethyl)benzyl (RS)-2-[2-[(1,1-dimethylethyl)pyrimidin-5-yl]-3,3-dimethylbutanoate (II). II at 500 ppm gave 50-79% mortality against *Blattella germanica* and 80-100% mortality against 9 addnl. pest species, e.g. *Tetranychus urticae*, *Nephotettix cincticeps*, and *Diabrotica balteata*. An emulsifiable concn. compn. contg. Sympersonic OP10 3.0, calcium dodecylbenzenesulfonate 2.0, and Aromasol H 94.0 wt. % was prepd.

L2 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1988:565740 CAPLUS
 DOCUMENT NUMBER: 109:165740
 ORIGINAL REFERENCE NO.: 109:27391a,27394a
 TITLE: Diacetone alcohol-comprising nonaqueous liquid ectoparasiticide pour-on formulation
 Metzner, Helmut; Steiner, Theodor; Mayer, Peter
 INVENTOR(S): Ciba-Geigy A.-G., Switz.
 PATENT ASSIGNEE(S): Eur. Pat. Appl., 22 pp.
 SOURCE: CODEN: EPXXDW
 DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 273862	A2	19880706	EP 1987-810705	19871130
EP 273862	A3	19880727		
EP 273862	B1	19910710		
R: AT, BE, CH, DE, ES, FR, GB, GR, IT, LI, LU, NL, SE				
AT 65013	T	19910715	AT 1987-810705	19871130
ES 2037740	T3	19930701	ES 1987-810705	19871130
IL 84701	A	19910415	IL 1987-84701	19871203
AU 8782112	A	19880609	AU 1987-82112	19871204
AU 603163	B2	19901108		
JP 63156716	A	19880629	JP 1987-307375	19871204
ZA 8709114	A	19880831	ZA 1987-9114	19871204
			CH 1986-4865	A 19861205

PRIORITY APPLN. INFO.:
 OTHER SOURCE(S): MARPAT 109:165740

AB The title formulation comprises an active ingredient(s), diacetone alc., and auxiliary ingredients. A formulation comprising 2% cypermethrin and 98% diacetone alc., applied at 2 mL/10 kg, totally controlled *Damalinea ovis* on sheep. An aqueous solution of 2-cyano-3-cyanoamino-3-methylthioacrylonitrile Na salt was treated with HCl, to give 2-chloro-4-amino-5-cyano-6-methylthiopyrimidine, which was suspended in acetonitrile and treated with cyclopropylamine, under refluxing, to give 2-cyclopropylamino-4-amino-5-cyano-6-methylthiopyrimidine. This was treated with NH₃ in an autoclave, at 150°, for 15-20 min, to give 2-cyclopropylamino-4,6-diamino-5-cyanopyrimidine.

10/546,138>

04/13/2008

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LOGOFF? (Y)/N/HOLD:Y

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

20.71

20.92

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE

TOTAL

ENTRY

SESSION

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-4.00

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